

Supplementary Material
A Novel High-Sensitivity Fluorescent Detection
Technology for Zearalenone Based on PER-Triggered
crRNA Conformational Change and CHA-Coordinated
Energy Supply

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1. Experimental section

1.1 Reagents and materials

Oligonucleotide, N,N,N',N-tetramethyl ethylenediamine (TEMED), ammonium persulfate (AP), 30% acrylamide solution (29:1), DEPC-treated water were purchased from Shanghai Sangong Biotechnology Co., Ltd. (Shanghai, China). $10 \times$ TBE Buffer, DNA Marker and GelRed were purchased from thermo fisher Co., Ltd (Beijing, China). Microcystin (MC-LR) were purchased from Macklin Inc. (Shanghai, China). Cas12a protein (Lba Cas12a) were purchased from New England Biolabs. Saxitoxin were purchased from Sichuan Hengcheng Zhiyuan Biotechnology Co., Ltd. Zearalenone(ZEN), Ochratoxin A(OTA), Dimethyl Sulfoxide(DMSO)were purchased from Shanghai Aladdin Biochemical Technology Co.,Ltd.Ochratoxin B(OTB),Aflatoxin B₁(ATB₁) were purchased from Sigma-Aldrich(Shanghai)Trading Co.,Ltd.

1.2 Apparatus.

All fluorescence profiles were determined by an Agilent Cary Eclipse spectrophotometer (Agilent, USA). Polypropylene gel electrophoresis (PAGE) was first performed using a DYCZ-24DN electrophoresis apparatus (LIUYI, Beijing, China), followed by shaking with $1 \times$ GelRed, WD-9405F decolorization shaker (LIUYI, Beijing, China). and finally, a Bio-Rad gel imaging system (Bio-Rad, USA) was used to analyze the samples.

2. Supporting table.

The DNA fragments shown in Table S1 were all purified by HPLC. The reagents used in the experiment were all of analytical purity.

Table S1. Nucleotide sequences used in this study

Oligonucleotides name	Sequence (5' to 3')	description
Apt	TTAATCATCTATCTATGGTACATTACTATCTGTAATGTGATATG	
T	TGTACCATAGATAGATGAGTTAATG	
YL	AATTTCTACTAAGTGTTTCATTCAT/i2OMeG/GCCGTTTTTTTTTCG GC/i2OMeC/ATGAATGAACACTTAGTAGAAATTATTTTTGGTTT CCCATTAACTCATCTATCTATGGTACAGTTAATGGGAAACCAA AAAAT	
Primer	CAAAAAAT	
HP1	AUGAAUGA ACACUUAGUAGAAAUU AUUUUUUGGUUUC UCAAAGGG GACCAAAAAAU AAUUUCUACUAAGUGU AGAUGCUCUGUCCUAACAGGCUA	
HP2	AGGTC CCCTTTGA GAAACCAAAAAAT AATTTCTACTAAGTGT TTTATCAAAGGG	
NTS	GGGGTCTTTAGCTTCTGTCCTAACAGGCTAAGT	
TS	ACTTAGCCTGTTAGGACAGAAGCTAAAGACCCC	
F-Q	FAM-TTATT-BHQ	